

SEP 30 2003

K032875

**510(k) Summary: Aloka Model SSD-5500 V6.0 Diagnostic Ultrasound System**

This summary statement complies with 21 CFR, section 807.92 as amended March 14, 1995.

This premarket notification has been submitted by Aloka Co., Ltd. and covers the Aloka SSD-5500 V6.0 diagnostic ultrasound system and transducers. The address is:

10 Fairfield Boulevard  
Wallingford, CT 06492  
(203) 269-5088

The contact person is: Richard J. Cehovsky, RA/QA Coordinator

The proprietary name is the Aloka SSD-5500 V6.0 diagnostic ultrasound system and transducers. The common name for this type of device is a diagnostic ultrasound system and transducers.

The item in this submission is covered under the following classification:

90 IYN	Ultrasonic Pulsed Doppler Imaging System	21 CFR 892.1550
90 ITX	Diagnostic Ultrasound Transducer	21 CFR 892.1570
90 IYO	Ultrasonic Pulsed Echo Imaging System.	21 CFR 892.1560

The above as stated in 21 CFR, part 892.1570, 1560 and 1550, has been classified as regulatory Class II.

The Aloka SSD-5500 V6.0 and its transducers are substantially equivalent to its predicates; the Aloka SSD-1700 and SSD-5500 Versions 3.1, 4.2 and 4.2.2 and its transducers.

The Aloka SSD-5500 V6.0 functions in the same manner as other diagnostic ultrasound devices. High frequency sound waves are transmitted into the body by a piezo-electric transducer. In the body, differences in the acoustic impedance of different tissues reflect a certain amount of the ultrasound energy back to the transducer, where it is transmitted via the probe cable to the system console and processed into an image. The Aloka SSD-5500 V6.0 transducer can also use the Doppler shift of sound reflected from moving tissues (blood) to detect and display flow.

The Aloka SSD-5500 V6.0, like other Aloka marketed diagnostic ultrasound systems and transducers are indicated for imaging body structures to aid in the diagnosis of disease or abnormality.

The Aloka SSD-5500 V6.0 diagnostic ultrasound system and transducers are similar in technological characteristics to Aloka's initial predicate ultrasound system: SSD-1700 (K963616) as well as Aloka's SSD-5500 Ver.3.1 (K992663), SSD-5500 Ver.4.2 (K002784/ K011457) and SSD-5500 Ver. 4.2.2 (K011315).

- The SSD-5500 V6.0 is indicated for the same diagnostic ultrasound applications to Aloka's ultrasound systems: SSD-1700 (K963616), SSD-5500- V3.1, V4.2 , V4.2.2.
- The SSD-5500 V6.0 has the same gray-scale and Doppler abilities to Aloka's ultrasound systems as mentioned above.

**510(k) Summary: Aloka Model SSD-5500 V6.0 Diagnostic Ultrasound System**

- The SSD-5500 V6.0 uses essentially the same technologies for imaging, Doppler functions and signal processing as the following products currently marketed by Aloka : SSD-1700 - (K963616) and SSD-5500- V3.1, V4.2 and V4.2.2 .
- The SSD-5500 V6.0 has the same method of use as the following products currently marketed by Aloka: SSD-1700 - (K963616) and SSD-5500-V3.1, V4.2 , & V4.2.2.
- The SSD-5500 V6.0 acoustic power output levels are below the maximum levels allowed by the FDA.
- The SSD-5500 V6.0 is subjected to the same Quality Assurance systems in development and production as other products currently marketed by Aloka such as the: SSD-1700 -(K963616) and SSD-5500- V3.1,V4.2, & V4.2.2.
- The patient contact materials used in the SSD-5500 V6.0 have been evaluated for safety via the same standards and methods as the above mentioned products marketed by Aloka. These materials have been found to be safe for the intended uses.
- The SSD-5500 V6.0 complies with electrical and physical safety standards as other products currently marketed by Aloka such as the: SSD-1700 - (K963616) and SSD-5500- V3.1, & V4.2, V4.2.2.
- Aloka Co., Ltd. Certifies that the SSD-5500 V6.0 complies with NEMA-UD2: 1992, AIUM 1994 "Acoustic Output Labeling Standard for Diagnostic Ultrasound Equipment", IEC-60601-1 (2001-09 Class A), UL 2601-1, 2<sup>nd</sup> edition (1997), Part 1, 2<sup>nd</sup> edition including Amendments 1&2 and ISO10993-1:1997. All testing has been completed and the results meet the requirements of the standards above.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

SEP 30 2003

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Donald James Sherratt  
Medical Stream Director  
Intertek Testing Services NA, Inc.  
70 Codman Hill Road  
BOXBOROUGH MA 01719

Re: K032875

Trade/Device Name: Aloka SSD-5500 V6.0 Diagnostic Ultrasound System

Regulation Number: 21 CFR §892.1550

Regulation Name: Ultrasonic pulsed doppler imaging system

Product Code: 90 IYN

Regulation Number: 21 CFR §892.1560

Regulation Name: Ultrasonic pulsed echo imaging system

Product Code: 90 IYO

Regulatory Class: II

Dated: September 11, 2003

Received: September 15, 2003

Dear Mr. Sherratt:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

This determination of substantial equivalence applies to the following transducers intended for use with the Diagnostic Ultrasound System, Model SSD-5500 V 6.0, as described in your premarket notification:

Transducer Model Number

ASU-67-10/7.5	UC140P-AL5	UCT140-AL-5	UST-533
UST-547	UST-672-5/7.5	UST-675P	UST-995-7.5
UST-987-7.5	ASU-1000C-3.5	ASU-1001	ASU-1002
ASU-1005	UST-2265-2	UST-5268P-5	UST-5271S-5

UST-5276-5	UST-5280-5	UST-5281-5	UST-5286-2.5
UST-5287-3.5	UST-5293-5	UST-5294-5	UST-5296
UST-5524-5	UST-5524-7.5	UST-5526L-7.5	UST-5531
UST-5534T-7.5	UST-5536-7.5	UST-5539-7.5	UST-5543
UST-5545	UST-5548	UST-5712	UST-5713T
UST-9101-7.5	UST-9102-3.5	UST-9104-5	UST-9114-3.5
UST-9115-5	UST-9118	UST-9119	UST-9120
UST-9121	UST-9126	UST-9128	UST-52101
UST-52104	UST-52108	UST-52109	UST-MC11-8731

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus permits your device to proceed to market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801, please contact the Office of Compliance at (301) 594-4591. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or at (301) 443-6597 or at its Internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Page 3 – Mr. Donald Sherratt

If you have any questions regarding the content of this letter, please contact Mr. Rodrigo Perez  
(301) 594-1212.

Sincerely yours,

  
Nancy C. Brogdon  
Director, Division of Reproductive,  
Abdominal and Radiological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosures

## 4.3.1

**Diagnostic Ultrasound Indications for Use Form**  
**SSD-5500 V6.0**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		P	P	P		P	P		See Below	
Abdominal		P	P	P		P	P		See Below	
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological		P	P	P		P	P		See Below	
Pediatric		P	P	P		P	P		See Below	
Small Organ (specify)		P	P	P		P	P		See Below	
Neonatal Cephalic		P	P	P		P	P		See Below	
Adult Cephalic		P	P	P		P	P		See Below	
Cardiac		P	P	P	P	P	P		See Below	
Transesophageal		P	P	P		P	P		See Below	
Transrectal		P	P	P		P	P		See Below	
Transvaginal		P	P	P		P	P		See Below	
Transurethral										
Intravascular										
Peripheral Vascular		P	P	P		P	P		See Below	
Laparoscopic		P	P	P		P	P		See Below	
Musculo-skeletal Conventional		P	P	P		P	P		See Below	
Musculo-skeletal Superficial		P	P	P		P	P		See Below	
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWDIntraoperative applications: include liver, pancreas and gall bladder. Small parts applications include breast, testes and thyroid.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*James C. Breyton*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
ASU-67-10/7.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal	P	P	P		P	P			See Below	
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UC140P-ALS**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal		P	P	P		P	P			See Below
Intraoperative (specify)		P	P	P		P	P			See Below
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal		P	P	P		P	P			See Below
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal										
Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD.

Intraoperative applications: liver, pancreas and gall bladder

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

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(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UCT140-AL-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
Ophthalmic										
Fetal										
Abdominal		P	P	P		P	P		See Below	
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal		P	P	P		P	P		See Below	
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative applications: include liver, pancreas and gall bladder.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-533**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)	E	E	E			E	E		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD,B/Bflow/PWD

Intraoperative applications: liver, pancreas and gall bladder.

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

KO32875

**Diagnostic Ultrasound Indications for Use Form**  
UST-547

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>	E	E	E			E	E		See Below	
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>	E	E	E			E	E		See Below	
<b>Neonatal Cephalic</b>	E	E	E			E	E		See Below	
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

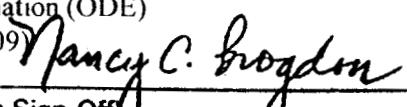
Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD.

Intraoperative applications: include liver, pancreas and gall bladder. Small Organ applications: breast, testes, thyroid

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)



(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number KO32875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-672-5/7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>		P	P	P		P	P		See Below	
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>		P	P	P		P	P		See Below	
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD.

Intraoperative applications: abdominal, bladder, pancreas and gall bladder.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-675P**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal	P	P	P			P	P		See Below	
Transvaginal	P	P	P			P	P		See Below	
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-995-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)	P	P	P		P	P			See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)	P	P	P		P	P			See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular	P	P	P		P	P			See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD.

Applications: Intraoperative: liver, pancreas and gall bladder. Small Organ: breast, testes, thyroid.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-987-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic		P	P	P		P	P		See Below	
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative applications: include liver, pancreas and gall bladder.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)

Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
ASU-1000C-3.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		P	P	P		P	P		See Below	
Abdominal		P	P	P		P	P		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy Chaydon*  
 \_\_\_\_\_  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
ASU-1001

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		P	P	P		P	P		See Below	
Abdominal		P	P	P		P	P		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

*KO32825*

**Diagnostic Ultrasound Indications for Use Form**  
ASU-1002

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
<b>Ophthalmic</b>										
<b>Fetal</b>		P	P	P		P	P		See Below	
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>		P	P	P		P	P		See Below	
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
ASU-1005

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		E	E	E		E	E		See Below	
Abdominal		E	E	E		E	E		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number K032825

**Diagnostic Ultrasound Indications for Use Form**  
**UST-2265-2**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>						P				See Below
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.10)

*Nancy C. Brogdon*  
(Division Sign-Off)

Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_

*K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5268P-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>		P	P	P		P	P		See Below	
<b>Intraoperative Neurological</b>		P	P	P		P	P		See Below	
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>		P	P	P		P	P		See Below	
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Neurological burr hole, Intraoperative: liver, pancreas, gall bladder

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_ X032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5271S-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>		P	P	P		P	P		See Below	
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>		P	P	P		P	P		See Below	
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

Nancy C Brogdon  
(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number K032815

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5276-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric	E	E	E			E	E		See Below	
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac	E	E	E			E	E		See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number KO32875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5280-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Opthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac		P	P	P		P	P		See Below	
Transesophageal		P	P	P		P	P		See Below	
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5281-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic	P	P	P		P	P			See Below	
Adult Cephalic										
Cardiac	P	P	P		P	P			See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Fancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
UST-5286-2.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac	P	P	P	P	P	P			See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal										
Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 Nancy C. Brogdon  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
UST-5287-3.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac		E	E	E		E	E		See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

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(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices  
41510(k) Number *K832875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5293-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>		P	P	P		P	P		See Below	
<b>Transesophageal</b>										
<b>Transrectal</b>			.							
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

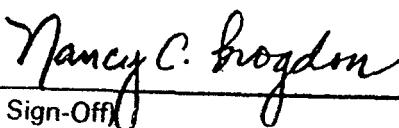
N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 Nancy C. Brogdon  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
UST-5294-5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic	P	P	P			P	P			See Below
Adult Cephalic										
Cardiac	P	P	P			P	P			See Below
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Neonatal

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5296**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic	P	P	P		P	P			See Below	
Adult Cephalic										
Cardiac	P	P	P		P	P			See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Neonatal

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5524-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)		P	P	P		P	P		See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular		P	P	P		P	P		See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Small Organ applications: breasts, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5524-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>		P	P	P		P	P		See Below	
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>		P	P	P		P	P		See Below	
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Small Organ applications: breasts, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5526L-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
<b>Opthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>		P	P	P		P	P		See Below	
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>		P	P	P		P	P		See Below	
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative: Liver, pancreas, gall bladder

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_

*KOB2875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5531**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative applications: include liver, pancreas and gall bladder.

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

*K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5534T-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>	P	P	P			P	P		See Below	
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>	P	P	P			P	P		See Below	
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>	P	P	P			P	P		See Below	
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Intraoperative- liver, pancreas and gall bladder. Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

**Diagnostic Ultrasound Indications for Use Form**  
UST-5536-7.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic		P	P	P		P	P		See Below	
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Intraoperative- liver, pancreas and gall bladder.

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number \_\_\_\_\_

KO32875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5539-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>		P	P	P		P	P		See Below	
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>		P	P	P		P	P		See Below	
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>		P	P	P		P	P		See Below	
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)

Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5543**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)		P	P	P		P	P			See Below
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular		P	P	P		P	P			See Below
Laparoscopic										
Musculo-skeletal Conventional		P	P	P		P	P			See Below
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number X032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5545**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)	P	P	P			P	P		See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular	P	P	P			P	P		See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial	P	P	P			P	P		See Below	
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*(Division Sign-Off)*

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number \_\_\_\_\_

*KQ32875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5548**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)	E	E	E			E	E		See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular	E	E	E			E	E		See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications- Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K032875

**Diagnostic Ultrasound Indications for Use Form**  
UST-5712

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)		P	P	P		P	P		See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular		P	P	P		P	P		See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications- Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Fancy C. Brogdon*  
(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number *KO32875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-5713T**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)		P	P	P		P	P		See Below	
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular		P	P	P		P	P		See Below	
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Intraoperative- liver, pancreas and gall bladder. Small parts: breast, testes and thyroid

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9101-7.5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>		P	P	P		P	P		See Below	
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>		P	P	P		P	P		See Below	
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_

*KO32875*

**Diagnostic Ultrasound Indications for Use Form**  
UST-9102-3.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal		E	E	E		E	E		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number

*X032876*

**Diagnostic Ultrasound Indications for Use Form**  
UST-9104-5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		P	P	P		P	P		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic		P	P	P		P	P		See Below	
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Intraoperative- liver, pancreas and gall bladder.

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*  
(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number *K032875*

**Diagnostic Ultrasound Indications for Use Form**  
UST-9114-3.5

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		P	P	P		P	P		See Below	
Abdominal		P	P	P		P	P		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9115-5**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Opthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>		P	P	P		P	P		See Below	
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>		P	P	P		P	P		See Below	
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number \_\_\_\_\_

*K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9118**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>		P	P	P		P	P		See Below	
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>		P	P	P		P	P		See Below	
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number \_\_\_\_\_ K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9119**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
<b>Ophthalmic</b>										
<b>Fetal</b>		P	P	P		P	P		See Below	
<b>Abdominal</b>		P	P	P		P	P		See Below	
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

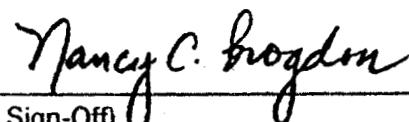
N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)



(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices

63 510(k) Number



**Diagnostic Ultrasound Indications for Use Form**  
**UST-9120**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>		E	E	E		E	E		See Below	
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>		E	E	E		E	E		See Below	
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Intraoperative- liver, pancreas, gall bladder

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

*K032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9121**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>		P	P	P		P	P			See Below
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>										
<b>Transesophageal</b>										
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal</b>										
<b>Conventional</b>										
<b>Musculo-skeletal</b>										
<b>Superficial</b>										
<b>Other</b>										

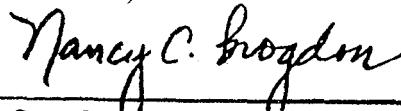
N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)



(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

K032875

**Diagnostic Ultrasound Indications for Use Form**  
UST-9126

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal		E	E	E		E	E		See Below	
Abdominal		E	E	E		E	E		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Abdominal and Gynecological, fetal  
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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

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Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number \_\_\_\_\_

K032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-9128**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal		E	E	E		E	E		See Below	
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric		E	E	E		E	E		See Below	
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

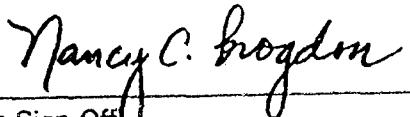
N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 Nancy C. Brogdon  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number X032875

**Diagnostic Ultrasound Indications for Use Form**  
**UST-52101**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac	E	E	E			E	E		See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number

*K030875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-52104**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

<b>Clinical Application</b>	<b>Modes of operation</b>									
	<b>A</b>	<b>B</b>	<b>M</b>	<b>PWD</b>	<b>CWD</b>	<b>Color Doppler</b>	<b>Amplitude Doppler</b>	<b>Color Velocity Imaging</b>	<b>Combined (specify)</b>	<b>Other (specify)</b>
<b>Ophthalmic</b>										
<b>Fetal</b>										
<b>Abdominal</b>										
<b>Intraoperative (specify)</b>										
<b>Intraoperative Neurological</b>										
<b>Pediatric</b>										
<b>Small Organ (specify)</b>										
<b>Neonatal Cephalic</b>										
<b>Adult Cephalic</b>										
<b>Cardiac</b>		N	N	N		N	N		See Below	
<b>Transesophageal</b>		N	N	N		N	N		See Below	
<b>Transrectal</b>										
<b>Transvaginal</b>										
<b>Transurethral</b>										
<b>Intravascular</b>										
<b>Peripheral Vascular</b>										
<b>Laparoscopic</b>										
<b>Musculo-skeletal Conventional</b>										
<b>Musculo-skeletal Superficial</b>										
<b>Other</b>										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number

*1032875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-52108**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic		E	E	E		E	E		See Below	
Adult Cephalic										
Cardiac		E	E	E		E	E		See Below	
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Applications: Neonatal

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,  
and Radiological Devices  
510(k) Number

*K032815*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-52109**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		E	E	E		E	E		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic		E	E	E		E	E		See Below	
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative applications: Liver, pancreas, gall bladder

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

(Division Sign-Off)

Division of Reproductive, Abdominal,

and Radiological Devices

510(k) Number

*KO32875*

**Diagnostic Ultrasound Indications for Use Form**  
**UST-MC11-8731**

Intended Use: Diagnostic ultrasound imaging or fluid flow analysis of the human body as follows:

Clinical Application	Modes of operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)		E	E	E		E	E		See Below	
Intraoperative Neurological										
Pediatric										
Small Organ (specify)										
Neonatal Cephalic	E	E	E		E	E			See Below	
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other										

N= new indication; P= previously cleared by FDA; E= added under Appendix E

Additional Comments: Mixed mode operation includes B/M, B/PWD, B/Bflow/PWD

Intraoperative applications: Liver, pancreas, gall bladder

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*Nancy C. Brogdon*

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Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number

*K032875*